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Dalmaprost 0,075 mg/ml raztopina za injiciranje za govedo, prašiče in konje

Ima dovoljenje za promet

- R-Cloprostenol sodium

Informacije o zdravilu

Ime zdravila:
Dalmaprost 0,075 mg/ml raztopina za injiciranje za govedo, prašiče in konje

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
krava
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
0.08 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **krava**
  - Meat and offal. no withdrawal period — withdrawal period is 0 days
  - Milk. no withdrawal period — withdrawal period is 0 hours
- **Pig (female)**
  - Meat and offal. 1 day
- **Horse (mare)**
  - Meat and offal. 2 day
  - Milk. no withdrawal period — withdrawal period is 0 hours

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Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

QG02AD90

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:

Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:
3/09/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Fatro S.p.A.

Pristojni organ:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Številka dovoljenja za promet z zdravilom:
DC/V/0680/001

Datum spremembe statusa dovoljenja za promet:
3/09/2019

Referenčna država članica:

Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Številka postopka:
ES/V/0305/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

slovenščina (PDF)
Objavljeno na dan: 13/04/2026
Prenesi

Navodilo za uporabo

slovenščina (PDF)
Objavljeno na dan: 13/04/2026
Prenesi

Označevanje

slovenščina (PDF)
Objavljeno na dan: 13/04/2026
Prenesi

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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